

Food and Drug Administration Rockville MD 20857

FILE COPY

February 8, 2002

Mr. Craig Patnode SterilMed, Inc. 11400 73rd Avenue North Minneapolis, MN 55369

Dear Mr. Patnode:

Your petition requesting the Food and Drug Administration to modify the 2/14/02 deadline for agency clearance of 510(k) for Class II reprocessed devices & PMA approval deadline for Class III reprocessed devices, was received by this office on 02/08/02. It was assigned docket number 02P-0061/CP1 and it was filed on 02/08/02. Please refer to this docket number in future correspondence on this subject with the Agency.

Please note that the acceptance of the petition for filing is a procedural matter in that it in no way reflects an agency decision on the substantive merits of the petition.

Sincerely,

Helen K. Hárris,

Dockets Management Branch

028-0061

ACKI